

Veterinary Feed Directive for Swine
ChlorMax®
(chlortetracycline)

Veterinarian: _____

Client: _____

Address: _____

Business or
Home Address: _____

Phone #: _____

Phone #: _____

FAX or email: (optional) _____

FAX or email: (optional) _____

Indications, Drug Level, and Duration of Use: (select one and specify additional required information)

- ☐ 1) Swine: Control of porcine proliferative enteropathies (ileitis) caused by *Lawsonia intracellularis* susceptible to chlortetracycline.

Drug level: _____ g/ton (approximately 400 g/ton to provide 10mg/lb BW/day)

Duration of use: _____ days (feed for not more than 14 days)

- ☐ 2) Swine: Treatment of bacterial enteritis caused by *Escherichia coli* and *Salmonella choleraesuis* and bacterial pneumonia caused by *Pasteurella multocida* susceptible to chlortetracycline.

Drug level: _____ g/ton (approximately 400 g/ton to provide 10mg/lb BW/day)

Duration of use: _____ days (feed for not more than 14 days)

USE OF FEED CONTAINING THIS VETERINARY FEED DIRECTIVE (VFD) DRUG IN A MANNER OTHER THAN AS DIRECTED ON THE LABELING (EXTRA-LABEL USE) IS NOT PERMITTED.

Approximate number of **Swine** to be treated: _____

Premises or Location of animals: _____

Special Instructions and/or other animal identifications:



Affirmation of Intent (for combination VFD drugs): check the appropriate box:

- ☐ This VFD only authorizes the use of the VFD drug(s) cited in this order and is not intended to authorize the use of such drug(s) in combination with any other animal drugs.
- ☐ This VFD authorizes the use of the VFD drug(s) cited in this order in the following FDA-approved, conditionally approved, or indexed combination(s) in medicated feed that contains the VFD drug(s) as a component.

	Drug(s) and Dose Range(s)	Specifications*
<input type="checkbox"/>	10 to 30 g/ton bacitracin methylenedisalicylate; (BMD®) [NADA 141-059]	Growing/finishing swine, pregnant sows This combination has a zero day withdrawal.
<input type="checkbox"/>	Other FDA-approved, conditionally approved, or indexed combination:	

*for complete information see the approved Type C medicated feed label

- ☐ This VFD authorizes the use of the VFD drug(s) cited in this order in any FDA-approved, conditionally approved, or indexed combination(s) in medicated feed that contains the VFD drug(s) as a component.

 **Warning:** Indication 1, 2: No withdrawal period required. 

Date of VFD Issuance: _____ (dd/mm/yyyy)

Date of VFD Expiration: _____ (dd/mm/yyyy)
(Cannot exceed 6 months after issuance)

Veterinarian's signature: _____

Color Z Original – Veterinarian

Color X Copy – Supplier

Color Y Copy – Client

All parties must retain a copy of this VFD for 2 years after issuance